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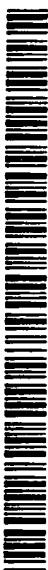
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(54) Title: A MOLD FOR NANO IMPRINTING

(57) Abstract: A metal mold for use in a nano-imprinting process comprises a firmly adhering monomolecular non-sticking layer. The layer was obtained by subjecting the mold to a reaction with a fluoroalkyl compound having a mercapto group. As a result of said reaction, the layer comprises an organic sulfide of said metal.

A MOLD FOR NANO IMPRINTINGTechnical field of the invention

The present invention relates to nano-imprinting in general and especially to a mold for nano-imprinting.

Background art

5 In the replication of nanostructures by hot embossing it is necessary to provide a non-sticking interface between the nanostructured master and the thermoplastic to be embossed. This allows demolding without degrading the replication fidelity. Adhesion during the mold release can cause damage to the replica. Further, residual 10 polymer structures can contaminate the surface of the mold. A prerequisite for a successful imprint process is that the mold used should be chemically and mechanically stable and also adhere poorly with polymers to avoid 15 sticking thereof to the mold.

R. W. Jaszewski, et al disclose in Microelectronic Engineering 35 (1997) 381-384 covering the surface of the mold with an ultra-thin anti-adhesive layer comprising a polytetrafluoroethylene (PTFE) film. The film is deposited 20 according to two kinds of techniques, i.e. plasma polymerization and ion sputtering from a plasma. Jaszewski et al allege in the above article that the quality of the film is degraded during the embossing process. Thus, it seems that the anti-adhesive layer according to Jaszewski 25 et al is not sufficiently stable.

Summary of the invention

It is a main object of the invention to provide a metal mold having improved anti-adhesive properties. This object is accomplished with a metal mold for use in a 30 nano-imprinting process, said mold comprising a firmly adhering monomolecular non-sticking layer, said layer being obtained by subjecting the mold to a reaction with a fluoroalkyl compound having a mercapto group, said

layer, as a result of said reaction, comprising a sulfide of said metal.

Other advantages and characteristics of the invention will be described below in more detail.

5 Detailed description of the invention

Before explaining the invention in greater detail a discussion will be presented concerning the imprint processes, to which the mold is applied. A non-limiting example is nano-imprint lithography, which is a technique 10 for mass production of nano structures. This includes two basic parts. The first part is a mold, which defines the nano structure on its surface, and the second part is pattern transfer from the mold into a polymer on various substrates. As to nano-imprint lithography reference is 15 made to US patent no. 5,772,905 to Chou and US patent no. 5,259,926 to Kuwabara et al. These patents are entirely incorporated herein by reference. The mold according to the invention can be used in other imprint processes as well.

20 In the present description "metal mold" refers to any mold for nano-imprinting that has a metal surface which is intended to contact the surface of a substrate to be embossed.

The term "nano-imprinting" should not be read as 25 referring only to submicron structures, i.e. structures with a size in the range of 1-1000 nm. A mold often includes both submicron structures and structures with a size of up to 100 micron, and larger. The present invention is applicable for molds comprising submicron structures and/or micron structures. The best effect of the 30 invention is observed with the submicron structures since these are relatively more sensitive to adhesion during mold release.

The present invention is based on the finding that a 35 very firmly adhering monomolecular non-sticking layer can be obtained on the mold by subjecting the latter to a reaction with a fluoroalkyl compound having a mercapto

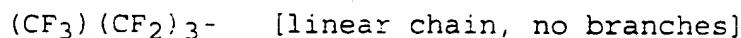
group. This compound includes two principal groups, R and X. The first group, called R, has non-sticking properties. The second group, X, which is bound to the R-group, binds the compound, denoted RX, to the mold surface to form a monomolecular non-sticking layer.

According to one aspect of the invention, the mold is made of metal, such as nickel, chromium, etc and thus the non-sticking layer is formed by reaction with a fluoroalkyl compound having a mercapto group. The result of the reaction is a very strong layer comprising an organic sulfide of nickel in case of a nickel mold.

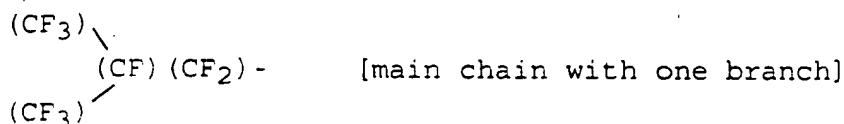
The R-group may include two subgroups; a first subgroup, called RA, that includes the non-sticking functionality, and a second subgroup, called RS, which serves as a "spacer".

The first subgroup, RA, is a fluoroalkyl-containing group. The simplified representation of the RA-group is $C_nF_mY_p$. C is carbon. The number of carbon atoms in RA is preferably n= 1-20, more preferred n= 1-10. F is fluorine. The number of fluorine atoms in RA is preferably m=1-30, more preferred m=3-20. Y may be hydrogen, oxygen, nitrogen, silicon or other atoms. In order to avoid any unwanted reactions with the surface of the mold Y is preferably hydrogen. The number of Y-atoms, p, is the number required to balance the RA-group with respect to its charge.

Each carbon atom in the RA-subgroup may be bound to 1-3 fluorine atoms. Preferably RA includes at least one perfluorinated carbon atom. Preferably the perfluorinated carbon atom is the terminal end carbon atom of the main chain of the RA-subgroup. RA may be linear or branched. For the non-sticking properties it is advantageous to have as many and as closely packed molecules of the RX-compound as possible bound to the surface of the mold. For reasons of close packing of the RX molecules the RA subgroup is preferably a linear chain, such as for example



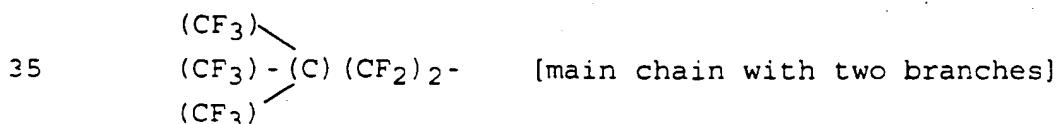
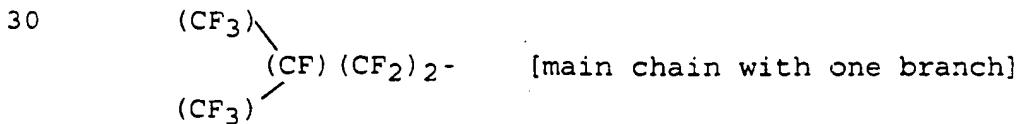
or a main chain with 1-3 short branches, such as for
5 example

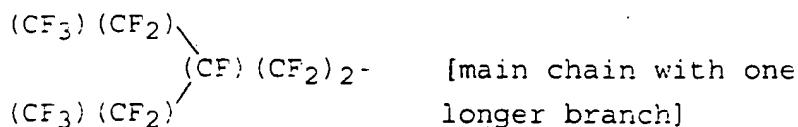


10

The branches, if any, preferably have a length of 1-3 carbon atoms each.

The surface of the metal mold for nano-imprinting has a certain nano structure. This structure could be 15 described as for example ridges, mountains and valleys in a nano scale. For some structures it may be difficult to obtain a sufficiently dense concentration of non-sticking functionality at the peaks of the mountains and at the corners of the ridges using a monomolecular layer of 20 linear molecules. Thus the terminal end of the RA main chain is preferably branched to make the "head" of the RX-compound larger to improve the density of the non-sticking functionality and thus the non-sticking properties at the peaks, edges and corners of the nano 25 structured surface. Preferably the branches as well as the main chain each include a perfluorinated carbon atom. Examples of such RA-subgroups with an enlarged head with non-sticking properties include:

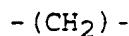




5 Another possibility, which is sometimes attractive for solving the above-mentioned difficulty, is to use very short main chains, for example R-groups with a total length of 2-3 carbon atoms, with no or short branches.

The non-sticking subgroup RA of the R-group should
 10 be situated at some distance from the mold surface to obtain the maximum non-sticking effect. The distance, represented as the number of carbon atoms being in line, from the X-group to the final and most distant carbon atom of the RA-subgroup is preferably 1-20 carbon atoms.
 15 Thus the total length of the main chain of the R-group is preferably 2-21 carbon atoms. If the RA-subgroup itself is not long enough a spacer subgroup RS may be used. The spacer subgroup RS is a hydrocarbon with the general formula C_qY_z . C is carbon and q is the number of carbon atoms. For linear RS-groups q is preferably 1-19. For branched RS-groups q is preferably 2-30. As discussed above the total distance from the X-group to the final carbon atom of the RA-subgroup is 1-20 carbon atoms. For
 20 a given RA-subgroup this condition results in a range for q. If the RA-subgroup itself has a sufficiently long carbon chain, then the RS-subgroup may be omitted. Y may be hydrogen, oxygen, nitrogen, silicon or other atoms.
 25 For reasons discussed above Y is preferably hydrogen.

The fluorine atoms of the RA-subgroup may influence
 30 the bond between the mercapto group and the metal. To avoid this it is preferable to use an RS-subgroup containing at least one hydrocarbon unit with the structure:



35

The unit $-(\text{CH}_2)-$ "isolates" the fluorine atoms of the RA-subgroup from the bond. Preferably an alkyl-group with at

least 2-3 units of this type should be used for best "isolation" of the RA-subgroup from the mercapto-group. The shortest RS-subgroup is thus preferably:

5 - (CH₂) (CH₂) -

For reasons of close packing of the RX-compound the RS-subgroup is preferably a linear group, such as for example

10

- (CH₂) (CH₂) (CH₂) - [linear chain, no branches]

or a group with short branches such as for example

15

- (CH₂) \ / (CH) (CH₂)₄ - [main chain with one branch]
- (CH₂) /

or

20

- (CH₂) \ / (CH) (CH₂)₄ (CH) / (CH₂) -
- (CH₂) / \ (CH₂) - [main chain with one branch at each end]

The branches, if any, preferably have a length of 1-3
25 carbon atoms each. Preferably the RS-subgroup is an alkyl-group. In the case of a branched RS-subgroup several RA-subgroups and/or several mercapto groups may be attached to the RS-subgroup. RA- and RS-subgroups containing double or triple bonds are possible but less
30 preferred since they may participate in unwanted addition reactions during heating of the mold.

The mold used in the nano-imprint process is a metal mold. The metal mold may be of two different types. The first type is a mold made from solid metal. The nano
35 structure is formed directly on the metal surface by e.g. an etching process known in the art. The second type is a mold comprising a base, e.g. made from silicon. The base

may be solid or made from several layers, and may be provided with a surface layer or film that is made from a metal. A preferred embodiment of this second type of mold is to first form the nano structure on a silicon surface, 5 by e.g. etching processes known in the art. Then a very thin layer of metal is applied on top of the silicon surface using e.g. thermic evaporation or electrochemical precipitation. The thickness of the metal layer depends on the nano structure and the required wear resistance of 10 the mold. In this way a mold is obtained having a metal surface that has a well defined nano structure and good wear properties.

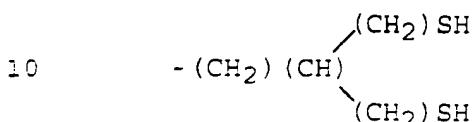
Several different metals, such as nickel, chromium, titanium, platinum, copper, silver, and gold, may be used 15 for the mold. The preferred metals are nickel, Ni, and chromium, Cr, due to their corrosion resistance, moderate price and high hardness. An important aspect is that the non-sticking layer on the metal mold should have very uniform properties. The binding process and the binding 20 energy of the monomolecular non-sticking layer will be different for different metals. The metal surface is thus preferably a rather pure metal, such as unalloyed nickel or unalloyed chromium. Alloys, such as stainless steel, and mixtures, such as a mixture of e.g. chromium and 25 nickel, are less preferred, since the properties of the monomolecular non-sticking layer bound to an alloy or a mixture will vary over the surface.

The X-group binds the RX-compound to the surface of the metal mold. The X-group includes at least one 30 mercapto group. One such mercapto group is the SH unit, wherein S denotes sulfur and H denotes hydrogen. The sulfur atom is bound directly to one of the carbon atoms of the R-group. If an RS-subgroup is included, then the sulfur of the SH unit is bound to one of the carbon atoms 35 in that subgroup. If no RS-subgroup is present then the sulfur is bound to one of the carbon atoms in the RA- subgroup. The X-group may contain more than one SH unit,

for example 2-4 SH units. In that case each SH-unit is bound to one carbon atom. The RS-subgroup bound to a X-group may, for example, have the following structure:

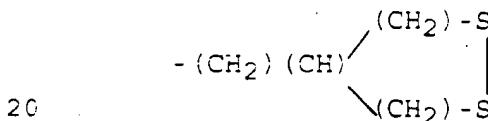
5 $-(\text{CH}_2)(\text{CH}_2)\text{SH}$ [RS-subgroup with X-group being
one SH-unit]

or



[branched RS-subgroup with X-group being two SH units]

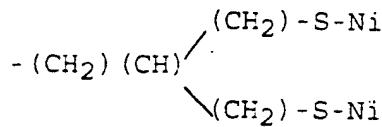
15 The SH units may, if they have a number of 2-4 in
each molecule of the RX-compound, form internal sulfur
bridges, that is an S-S unit, such as for example:



[branched RS-subgroup with X-group being a S-S unit]

25 The S-S unit is also included under the definition
mercapto group, since the S-S unit reacts with metal
surfaces in exactly the same way as the SH units and is
basically just another state of SH units.

30 The mercapto groups react readily with the metals to
form metal sulfides, such as nickel sulfide or chromium
sulfide. An RX-compound, where X includes two or more SH
units or at least one S-S unit, provides a stronger
attachment of the RX-compound to the surface. This is
preferred in cases where each metal sulfide bond is weak
or when an extra strong attachment is required. An
example of such an attachment, using a molecule with two
35 SH units or a molecule with a sulfur bridge, that is an
S-S unit, is shown below:

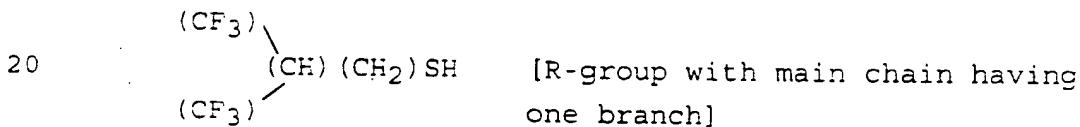


5 [branched RS-subgroup with X-group being two SH units or one S-S unit after attachment to a nickel surface]

10 For extremely small nano structures very small RX-compounds may be preferable. The main chain of the R-group may then be as short as 2-3 carbon atoms. Examples 15 of such RX-compounds, with R-groups having in total only 2-5 carbon atoms include:



and



25 The conditions during the reaction between the metal surface and the RX-compound are such that the compound is in a gaseous phase or in a liquid phase while the mold is 30 in a solid phase.

For reactions with the RX-compound in a gaseous phase it is preferred to clean the mold and place it in a vacuum oven under low pressure (about 10-200 mbar, preferably about 50-100 mbar) and at a temperature of about 50-200°C, preferably about 70-120°C. The fluoroalkyl compound having a mercapto group (the RX-compound) is then introduced into the oven. The mercapto group reacts with the metal surface forming a metal sulfide. When the reaction is completed the mold is withdrawn from the oven, cleaned, allowed to cool down 35 and is then ready for use.

For reactions with the RX-compound in a liquid phase the mold is preferably cleaned in a solvent being adapted to the RX-compound. Examples of such a solvent include alkanes like pentane, hexane, heptane, octane etc. A 5 solution is prepared by dissolving the RX-compound in a solvent, preferably the same solvent as was used for cleaning the mold. The concentration of the RX-compound in the solution is not critical, but often 1-10% is convenient. The temperature of the solution is not 10 critical, but for practical reasons room temperature is often convenient. The cleaned mold is then dipped into an agitated container containing the solution. The mercapto group reacts with the metal surface forming a metal sulfide. The mold is then removed from the container and 15 is cleaned by dipping it into one or several consecutive baths each preferably containing the same solvent as was used for the solution. The mold is finally dried, either by heating it sufficiently to evaporate the solvent or by flushing it with nitrogen gas.

20 The choice of RX-compounds of the formula above is not critical according to the invention and one skilled in the art can easily, without extensive experimentation, find the most appropriate one in each case. By this choice possibility the layer can be tailored in view of 25 factors material to the actual use of the mold, which is a great advantage.

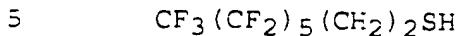
Best mode of performing the invention.

An embodiment of the invention, which at present is judged to be the best mode of performing the invention, 30 will be described below with reference to a working example.

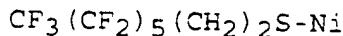
Example 1

A silicon mold was etched to define a certain nano structure. The surface of the mold was then coated with a 35 thin layer of nickel. The mold was cleaned in heptane.

A fluoroalkyl compound having a mercapto group, which compound was 1H, 1H, 2H, 2H-perfluorooctanethiol and had the structure:



was dissolved in heptane to obtain a 2% solution. The temperature of the solution was about 20°C. The cleaned metal mold was put into an agitated beaker containing the
10 solution. The mold was now provided with a mono-molecular non-sticking layer with the structure:



15 The mold was taken up from the beaker and was cleaned by dipping it in each of three consecutive beakers each containing heptane. The mold was then heated to 70°C whereby the heptane was evaporated. The mold was tested in a nano-imprint lithography process and it was found
20 that the mold withstood more than 50 imprint sequences without any detectable degradation.

The example above was repeated, except that the following fluoroalkyl compounds having a mercapto group were used:

25

Table 1: Additionally tested compounds

<u>R-group</u>		<u>X-group</u>	<u>Metal</u>
<u>RA-subgroup</u>	<u>RS-subgroup</u>		
$(\text{CF}_3)-$	$-(\text{CH}_2)_2-$	-SH	Ni
$(\text{CF}_3)_2(\text{CF})-$	$-(\text{CH}_2)(\text{CH})((\text{CH}_2)-)_2$	-S-S-	Cr
$(\text{CF}_3)(\text{CF}_2)_3-$	$-(\text{CH}_2)_2-$	-SH	Ni
$(\text{CF}_3)(\text{CF}_2)_7-$	$-(\text{CH}_2)_2-$	-SH	Cr

Similar results were obtained upon testing.

Example 2

30 The same RX-compound as described in example 1 was applied to a mold similar to that used in example 1. The

mold was cleaned in heptane and then put into an oven that had a temperature of 110°C, a nitrogen atmosphere and a pressure of 75 mbar. Pure RX-compound was injected into the oven. The mold, now provided with a mono-
5 molecular non-sticking layer with a structure similar to that described in example 1, was taken out of the oven and was cleaned by dipping it in each of three consecutive beakers each containing heptane. Finally, the mold was dried by heating it to 70°C. Similar results as in
10 example 1 were obtained upon testing.

CLAIMS

1. A metal mold for use in a nano-imprinting process comprising a firmly adhering monomolecular non-sticking layer, said layer being obtained by subjecting the mold to a reaction with a fluoroalkyl compound having a mercapto group, said layer, as a result of said reaction, comprising a sulfide of said metal.
- 10 2. A metal mold as claimed in claim 1, wherein the mold is made of solid metal or has a metal surface.
3. A metal mold as claimed in claim 2, wherein the metal mold comprises a thin layer of metal applied on a base material.
- 15 4. A metal mold as claimed in any of claims 1-3, wherein the metal is nickel or chromium.
5. A metal mold as claimed in any of claims 1-4, wherein the metal is unalloyed.
- 20 6. A metal mold as claimed in any of claims 1-5, wherein the fluoroalkyl compound having a mercapto group is denoted RX where;
R is a group with non-sticking properties, and
X includes at least one mercapto group,
wherein R is a linear fluoroalkyl group, which has a
25 first end and a second end and which has no or only short
branches, and wherein R at the first end includes at
least one fluorinated carbon atom and at the second end
is bound to X.
7. A metal mold as claimed in claim 6, wherein at
30 least one of said at least one fluorinated carbon atom is
a perfluorinated carbon atom.
8. A metal mold as claimed in any of claims 6-7, wherein R is a linear main chain, wherein the first end
of R includes 1-3 branches, each with a length of 1-3
35 carbon atoms, and wherein each of the branches and the
first end of R includes a perfluorinated carbon atom.

9. A metal mold as claimed in any of claims 6-7,
wherein the main chain of R has a total length of 2-3
carbon atoms.
10. A metal mold as claimed in any of claims 1-9,
5 wherein the nano-imprinting process is nano-imprint
lithography.
11. A metal mold as claimed in any of claims 1-10,
wherein during said reaction the fluoroalkyl compound is
in a gaseous phase.
- 10 12. A metal mold as claimed in any of claims 1-10,
wherein during said reaction the fluoroalkyl compound is
in a liquid phase or is dissolved in a solvent.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 01/00087

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: G03F 7/00, B41M 1/06, B81C 1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: B41M, B81C, G03F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

QUESTEL: EDOC, WPIL, JAPIO, DIALOG: DIALINDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5512131 A (AMIT KUMAR ET AL), 30 April 1996 (30.04.96), column 1, line 6 - line 24; column 8, line 29 - line 58; column 12, line 46 - line 52, figure 1 --	1-3,10-11
A	US 5358604 A (CHARLES W.C. LIN ET AL), 25 October 1994 (25.10.94), column 7, line 45 - line 61, figures 3,4 -- -----	1-12

<input type="checkbox"/>	Further documents are listed in the continuation of Box C.	<input checked="" type="checkbox"/> See patent family annex.
* Special categories of cited documents		

"A"	document defining the general state of the art which is not considered to be of particular relevance	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E"	earlier application or patent but published on or after the international filing date	"X"	document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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"O"	document referring to an oral disclosure, use, exhibition or other means	"&"	document member of the same patent family
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Date of the actual completion of the international search	Date of mailing of the international search report
11 April 2001	18-04-2001

Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. + 46 8 666 02 86	Authorized officer Bengt Christensson/MP Telephone No. + 46 8 782 25 00
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INTERNATIONAL SEARCH REPORT

Information on patent family members

25/02/01

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PCT/SE 01/00087

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5512131 A	30/04/96	US 5900160 A	04/05/99
US 5358604 A	25/10/94	NONE	